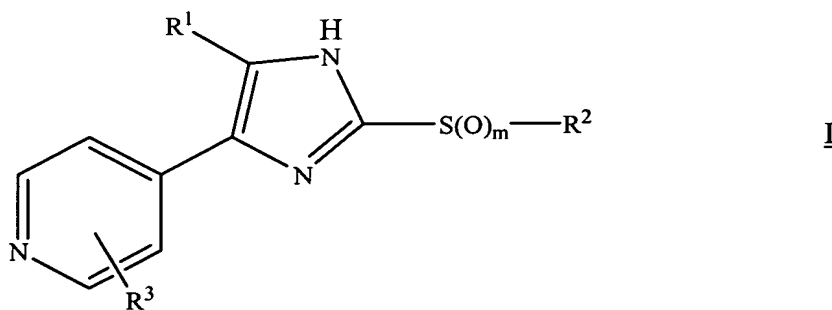


IN THE CLAIMS

Please amend the claims as follows:

Claim 1 (Currently Amended): A 2-thio-substituted imidazole derivative of the formula I:

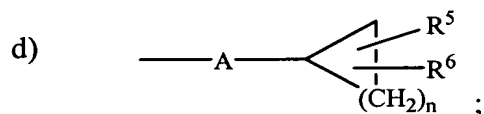
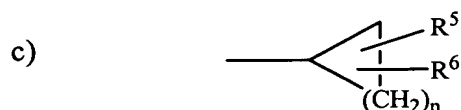


wherein ~~in which~~

R^1 is C_1 - C_6 -alkyl, C_3 - C_7 -cycloalkyl or aryl which is unsubstituted or substituted by a halogen atom, by C_1 - C_6 -alkyl or by halo- C_1 - C_6 -alkyl;

R^2 is selected from the group consisting of:

- a) aryl- C_1 - C_4 -alkyl, where the aryl radical may have one, two or three substituents, independently of one another, selected from the group consisting of C_1 - C_6 -alkyl, C_1 - C_6 -alkoxy, halogen, C_1 - C_6 -alkylsulfanyl, C_1 - C_6 -alkylsulfinyl, C_1 - C_6 -alkylsulfonyl and hydroxyl, and
- b) C_1 - C_6 -alkyl which is unsubstituted or substituted by CN or halogen[[:]],



R^3 is selected from the group consisting of:

- a) $NR^4R^{10}[[;]]_1$
- b) $NR^7COR^{10}[[;]]_1$
- c) $NR^7COOR^{10}[[;]]_1$
- d) $NR^7CONR^7R^{10}[[;]]_1$
- e) $NR^7CONR^7COR^{10}[[;]]_1$
- f) $OR^{10}[[;]]_1$
- g) $S(O)_mR^{10}_1$
- h) halogen[[;]]₁
- i) $OH[[;]]_1$
- j) N_3
- k) NH_2 , and
- l) $SH[[;]]_1$, and

wherein ~~where~~ R^3 is not OH, halogen, C₁-C₆-alkylthio or C₁-C₆-alkoxy, if R^2 is phenyl-C₁-C₄-alkyl₁ and the phenyl radical has a C₁-C₆-alkylsulfanyl, C₁-C₆-alkylsulfinyl or C₁-C₆-alkylsulfonyl substituent[[;]]₁

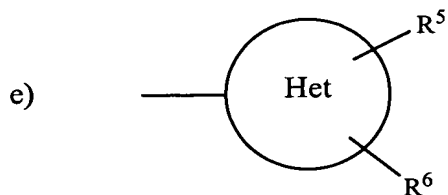
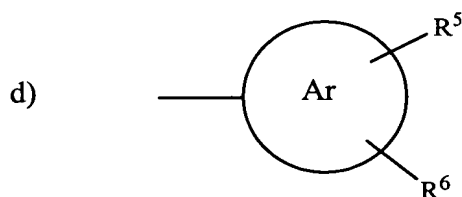
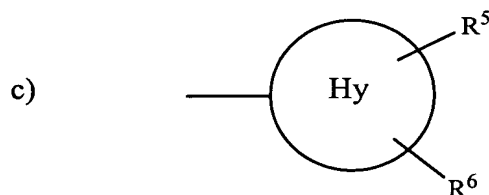
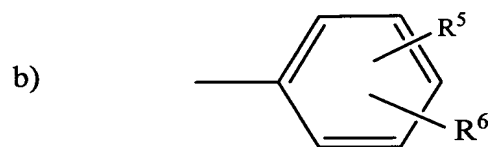
R^4 is H or a physiologically cleavable group,

R^5 and R^6 , which may be identical or different, are H, halogen, OH, C_1 - C_6 -alkoxy, C_1 - C_6 -alkyl, halo- C_1 - C_6 -alkyl, C_1 - C_6 -alkylsulfanyl, NH_2 , C_1 - C_6 -alkylamino or di- C_1 - C_6 -alkylamino[[:]],

R^7 is R^4 , C_1 - C_6 -alkyl or benzyl[[:]],

R^{10} is selected from the group consisting of ~~has one of the meanings below:~~

a) $A - B_2$



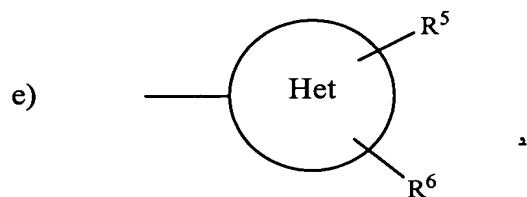
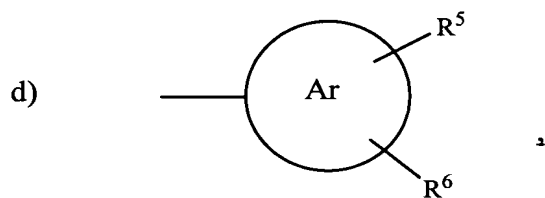
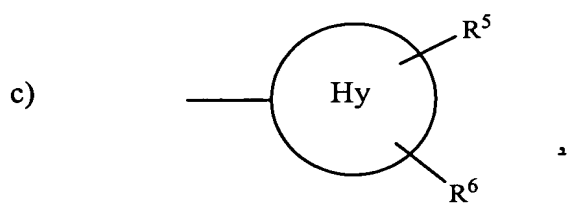
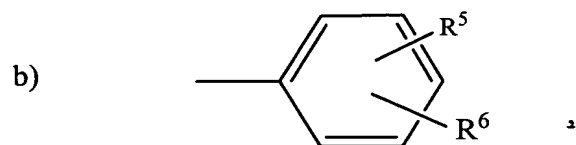
f) C_1 - C_6 -alkyl which is substituted by 2 or 3 phenyl groups[[:]], and

g) trifluoromethyl, Trifluoromethyl

A is straight-chain or branched C₁-C₆-alkylene, C₂-C₆-alkenylene or C₃-alkynylene[[:]],

B is selected from the group consisting of:

a) H,



f) OC₁-C₆-alkyl[[:]],

g) NR¹¹R¹²[[:]],

h) OH[[:]],

i) halogen[[:]], and

j) C₁-C₆-alkylsulfanyl,

R¹¹ and R¹², which may be identical or different, are H, C₁-C₆-alkyl or phenyl[[:]].

Hy is a 3- to 10-membered, non-aromatic mono-, bi- or tricyclic carbocycle, which may or may not be fused with a benzene ring[[:]],

Ar is a 5- or 6-membered aromatic heterocycle, which has 1, 2 or 3 hetero-atoms, independently of one another, selected from the group consisting of O, S and N, and which may or may not be fused with a benzene ring[[:]],

Het is a 5- or 6-membered, non-aromatic heterocycle, which has 1, 2 or 3 heteroatoms, independently of one another, selected from the group consisting of O, S and N, which may or may not be fused with a benzene ring, and which may or may not be bridged bicyclically or tricyclically;

m is ~~0-1~~ 0, 1 or 2;

n is 1, 2, 3, 4 or 5;

and the tautomers, optical isomers and physiologically acceptable salts thereof.

Claim 2 (Currently Amended): The imidazole derivative compound as claimed in claim 1 of the formula I, wherein ~~in which~~

R¹ is C₁-C₆-alkyl, C₃-C₇-cycloalkyl or aryl which may or may not be substituted by a halogen atom;

R² is selected from the group consisting of:

a) aryl-C₁-C₄-alkyl, where the aryl radical may have one, two or three substituents, independently of one another, selected from the group consisting of C₁-C₆-alkyl, C₁-C₆-alkoxy, halogen, C₁-C₆-alkylsulfanyl, C₁-C₆-alkylsulfinyl, C₁-C₆-alkylsulfonyl and hydroxyl, ~~and~~

b) C₁-C₆-alkyl which may or may not be substituted by CN; and

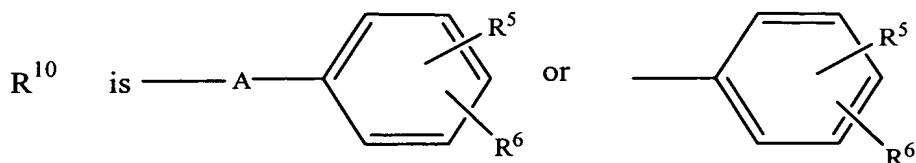
c) C₃-C₇-cycloalkyl;

R^3 is selected from the group consisting of:

- a) NR^4R^{10} ,
- b) NR^7COR^{10} , NR^7COR^{10} ,
- c) halogen and,
- d) C_1 - C_6 -alkoxy, and
- e) C_1 - C_6 -alkylthio,

where wherein R^3 is not OH, halogen, C_1 - C_6 -alkylthio or C_1 - C_6 -alkoxy, if R^2 is phenyl- C_1 - C_4 -alkyl, and the phenyl radical has a C_1 - C_6 -alkylsulfanyl, C_1 - C_6 -alkylsulfinyl or C_1 - C_6 -alkylsulfonyl substituent;

R^4 is $H[[:]]$,



or, if R^3 is NR^7COR^{10} , NR^7COR^{10} , is R^8 ,

R^5 and R^6 , which may be identical or different, are H, halogen, OH, C_1 - C_6 -alkoxy, C_1 - C_6 -alkyl[[:]],

R^7 is H, C_1 - C_6 -alkyl or benzyl[[:]],

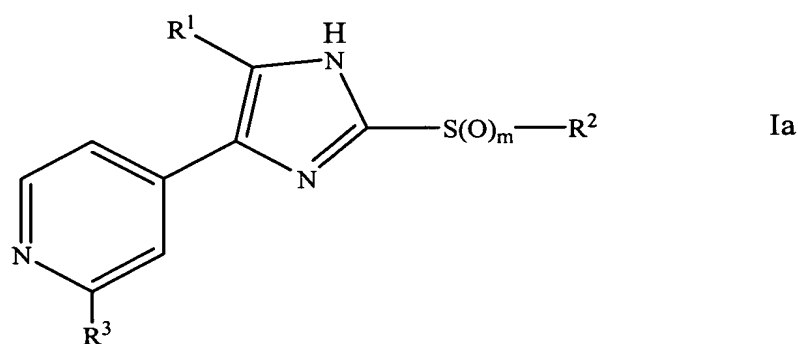
R^8 is C_1 - C_4 -alkyl, C_3 - C_6 -cycloalkyl or phenyl, where the phenyl group may have one or two substituents, independently of one another, selected from the group consisting of C_1 - C_4 -alkyl, C_1 - C_4 -alkoxy and halogen[[:]],

A is straight-chain or branched C₁-C₆-alkylene, C₂-C₆-alkenylene or C₃-alkynylene;
and

m is ~~0-1~~ 0, 1 or 2

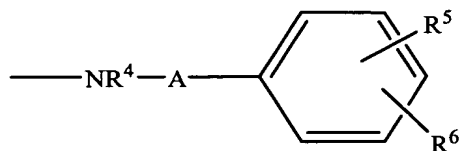
or a tautomer, an optic isomer or a physiologically acceptable salt thereof.

Claim 3 (Currently Amended): The imidazole derivative compound as claimed in claim 1 ~~or 2~~, wherein formula I of the is represented by the structure for formula Ia:



in which R¹, R², R³ and m are as defined for the structure of formula I in claim 1.

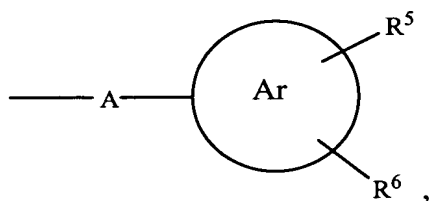
Claim 4 (Currently Amended): The imidazole derivative compound as claimed in claim 1 ~~or 2~~, wherein where R³ is



where A, R⁵ and R⁶ are as defined for the structure of formula I in claim 1.

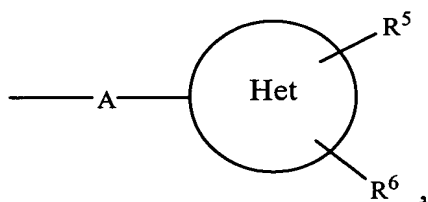
Claim 5 (Currently Amended): The imidazole derivative compound as claimed in claim 1, wherein, in of the formula I, in which R¹⁰ is selected from the group consisting of one of the radicals below:

a)



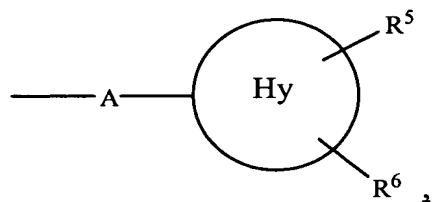
where Ar is a 5- or 6-membered aromatic heterocycle, which has a heteroatom selected from the group consisting of N, O and S; A is C₁-C₃-alkylene, and may be substituted by a phenyl radical, and R⁵ and R⁶ are H;

b)



where Het is a 5- or 6-membered non-aromatic heterocycle, which has an O or N heteroatom; A is C₁-C₃-alkylene, and R⁵ and R⁶ are H;

c)



where A is C₁-C₆-alkylene; R⁵ and R⁶ are H, and Hy is cyclopentyl or cyclohexyl;

d) cyclopentyl or cyclohexyl;

- e) phenyl-C₁-C₆-alkyl, where the alkyl radical may have an additional phenyl substituent; and
- f) C₂-C₆-alkenyl which is substituted by phenyl.

Claim 6 (Currently Amended): The imidazole derivative compound as claimed in claim 1, ~~wherein, in~~ of the formula I, ~~in which~~ R³ is A-B, and B is selected from the group consisting of NR¹¹R¹², OC₁-C₆-alkyl and OH, and A, R¹¹ and R¹² are as defined for formula I ~~in claim 1.~~

Claim 7 (Currently Amended): The imidazole derivative compound as claimed in claim 1, ~~wherein, in~~ of the formula I, ~~in which~~ R³ is NR⁷COR⁸, where R⁸ is selected from the group consisting of -O-C₁-C₄-alkylphenyl, phenyl and C₂-C₆-alkenyl which is substituted by phenyl.

Claim 8 (Currently Amended): The imidazole derivative compound as claimed in claim 1, wherein ~~any of the preceding claims where~~ A is C₁-C₂-alkylene.

Claim 9 (Currently Amended): The imidazole derivative compound as claimed in claim 1, wherein ~~any of the preceding claims where~~ A is ethylidene.

Claim 10 (Currently Amended): The imidazole derivative compound as claimed in claim 1, wherein ~~any of the preceding claims where~~ R⁵ and R⁶ are H.

Claim 11 (Currently Amended): The imidazole derivative compound as claimed in claim 1, wherein any of the preceding claims where R¹ is halogen-substituted phenyl, CF₃-substituted phenyl or C₁-C₆-alkyl-substituted phenyl.

Claim 12 (Currently Amended): The imidazole derivative compound as claimed in claim 1, wherein any of the preceding claims where R² is benzyl or C₁-C₆-alkyl.

Claim 13 (Currently Amended): A pharmaceutical composition, comprising the imidazole derivative at least one compound as claimed in claim 1, and if appropriate together with one or more pharmaceutically acceptable carriers and/or additives.

Claim 14 (Canceled)

Claim 15 (Currently Amended): A method for treating disorders associated with a disturbed immune system, which comprises, administering, to a person in need thereof, the imidazole derivative, characterized in that an amount of a compound of the formula I as claimed in claim 1, any of claims 1-12 in an amount sufficient to have immunomodulating action and/or to inhibit the release of cytokine ~~is administered to a person in need of such a~~ treatment.